

## **EXHIBIT C2**

2d ET Sm. JSMEL28 c Titration  
 of IETD

- Extracts made c 2M RA/DC pH 8.0, 20mM DTT, 100mM  
 - ET added 7pm, Inhib added 7pm (4hr later)  
 - Cells harvested @ ~1:30 pm Day 2 (~41 hours)  
 42.5 hr

PM	0.25 (4x)	0.25 (2x)	2.5	1x
2N	.264	1.9	1.32	0.53
2E	.346	2.8	0.89	0.36
2N/1m	.290	2.2	0.46	1.14
2E/1m	.326	2.5	1.0	0.4
2N/5m	.308	2.4	1.04	
2E/5m	.349	2.8	0.89	
2N/10m	.261	1.9	1.32	
2E/10m	.345	2.8	0.89	

Gel ① E CAT 2.5

2N 2E 2N/1 2E/1 2N/5 2E/5 2N/10 2E/10

Gel ① CAT 1.25 970 2.5

(M) 2N 2E 2N/5 2E/5 (M) 2N 2E 2N/5 2E/5

1.25 1.95  
 Cat 0.25 (4x) 8/7  
 2N .374 1.6  
 2E .482 2.2

## Addendum:

- 1nM IETD inhibits GCAD downreg and p120 downreg
- 1nM IETD suppresses  $\beta$ CAT in both stimulated samples and unstimulated controls

concl:

- 1nM IETD is sufficient to inhibit E-cad downreg
- higher concentrations may take out factors which participate in response
- ~~is~~ difficult to say if  $\beta$ CAT downreg is required for GCAD since IETD suppresses  $\beta$ CAT in unstimulated samples
- $\downarrow$  p120 may be a critical factor.

# Results

## ① ECAD

- Cell somewhat overloaded
- IETD inhibits GCAD downregulation @ 1nM but not @ 5nM and 10nM concentrations
- ? Toxicity of ECAD in 2E/IETD 1nM sample
- more potent ET-1 induced downregulation @ increasing concentrations of inhibitor

## Cell ②

### PRO

- SHIFT APPRECIATED! in ET-1 Amended sample
- 5nM IETD suppresses PRO levels @ baseline

- GCAD - 5nM IETD inhibits GCAD downregulation
- All 2E sample on day 1 so not sure if inhibition present

## ↳ Repeating ECAD

	18	<u>1nM</u>		PRO 2.5nM		<u>1nM</u>				
①	M	2N	2E	2N	2E	M	2N	2E	2N	2E
		5.3	3.6	4.5	4		13.3	9	11.3	10

	GCAD 18	<u>1nM</u>			<u>ext. 2.5nM</u>					
(2)	M	2N	2E	2N	2E	<del>2N</del>	2N	2E	4E	(M)
		5.3	3.6	4.5	4		7	8.3	7.8	